



Please amend the above-referenced application as follows:

✓  
In the specification:

Please replace Table 4, at page 43 of the specification with the following Table 4:

Table 4: Apparent kinetic rate and affinity constants of D2E7 and biotinylated rhTNF

C<sup>1</sup>

Experiment	$K_a$ ( $M^{-1}, s^{-1}$ )	$K_d$ ( $s^{-1}$ )	$K_d$ (M)
1	$1.33 \times 10^5$	$9.58 \times 10^{-5}$	$7.20 \times 10^{-10}$
2	$1.05 \times 10^5$	$9.26 \times 10^{-5}$	$8.82 \times 10^{-10}$
3	$3.36 \times 10^5$	$7.60 \times 10^{-5}$	$2.26 \times 10^{-10}$
Average	$1.91 \pm 1.26 \times 10^5$	$8.81 \pm 1.06 \times 10^{-5}$	$6.09 \pm 3.42 \times 10^{-10}$

C<sup>2</sup> Please insert drawings 1 to 9, filed herewith, after the last page of the specification.

In the claims:

Please cancel claim 1 without prejudice.

Please add the following new claims:

C<sup>3</sup>

74. (New) A pharmaceutical composition for treating a disorder in which TNF $\alpha$  activity is detrimental comprising an isolated human antibody, or an antigen-binding portion thereof, that dissociates from human TNF $\alpha$  with a  $K_d$  of  $1 \times 10^{-8}$  M or less and a  $K_{off}$  rate constant of  $1 \times 10^{-3} s^{-1}$  or less, both determined by surface plasmon resonance, and neutralizes human TNF $\alpha$  cytotoxicity in a standard *in vitro* L929 assay with an  $IC_{50}$  of  $1 \times 10^{-7}$  M or less, and at least one additional therapeutic agent.